

(19)



Europäisches Patentamt
European Patent Office
Office européen des brevets

(11) Publication number:

**0 314 078
A1**

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: **88117771.1**

(51) Int. Cl.4: **A61N 1/362**

(22) Date of filing: **25.10.88**

(30) Priority: **26.10.87 US 112861**

(43) Date of publication of application:
03.05.89 Bulletin 89/18

(84) Designated Contracting States:
DE FR GB IT NL SE

(71) Applicant: **MEDTRONIC, INC.**
7000 Central Avenue N.E.
Minneapolis Minnesota 55432(US)

(72) Inventor: **Mehra, Rahul**
901 Pine Street West
Stillwater, MN 55082(US)

(74) Representative: **Schwan, Gerhard, Dipl.-Ing.**
Elfenstrasse 32
D-8000 München 83(DE)

(54) **Ramped waveform non-invasive pacemaker.**

(57) A non-invasive electrical stimulator for pacing the heart. The stimulator generates a ramped current waveform which reduces muscle stimulation and pain associated with external stimulation of the heart. Each pulse (20) of the waveform has a current amplitude which rises gradually from 0 to a maximum current amplitude over a predetermined rise time (16) extending over at least about one-half of the duration of said pulse (20).

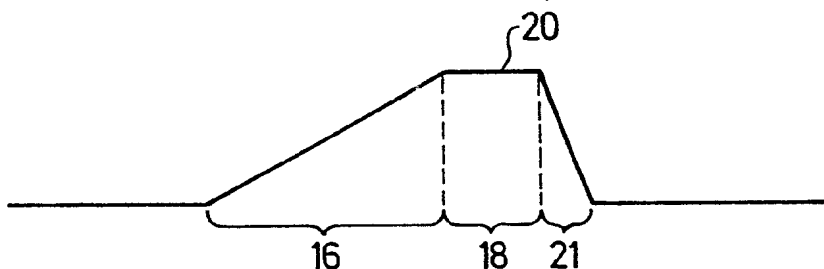


FIG. 2

EP 0 314 078 A1

NON-INVASIVE PACEMAKER

This invention relates generally to cardiac pacing and in particular to external, on-invasive cardiac pacing.

External non-invasive cardiac pacing is a well established technique which can provide effective heart pacing on a emergency basis. Traditionally, the pain and undesirable stimulation of skeletal muscles associated with non-invasive pacing have limited the acceptance of this technique.

Recent external pacemaker designs, such as that disclosed in US-A-4 349 030, have addressed the problem of pain associated with external non-invasive pacing by providing constant current pacing pulses of extended duration. Typically, these pulses are 20 to 40 ms in length and maintain a relatively constant current level throughout the duration of the pulse.

An object basic to this invention is to provide for a non-invasive pacemaker which is adapted to reduce pain and stinging associated with transthoracic pacing while reliably pacing the heart.

In conformity with the present invention a non-invasive cardiac pacemaker comprises:
first and second skin contacting electrode means for electrically coupling to a patient;
first timing means for repetitively timing an escape interval; and pulse generator means, coupled to and responsive to said first timing means and coupled to said first and second skin contacting electrode means, said pulse generator means generating pacing pulses for application to said patient via said first and second electrode means, said pacing pulses extending for a duration of at least about 20 ms, said pacing pulses having a current amplitude which rises gradually from 0 to a maximum current amplitude over a predetermined rise time of at least about 20 ms, said predetermined rise time extending over at least about one-half of the duration of said pacing pulse.

The improved external non-invasive cardiac pacemaker provided by the present invention can be tolerated by patients for an extended period of time. In particular, the present invention reduces the intensity of muscle stimulation associated with external non-invasive cardiac pacing by providing a ramped current pulse. In most patients, the use of a ramped current pulse as opposed to a constant current pulse significantly reduces skeletal muscle stimulation as compared to rectangular current pulses of the same amplitude and duration. In addition, use of a ramped waveform instead of a constant current waveform is believed to improve most patients' ability to tolerate extended periods of non-invasive pacing. Animal testing has revealed that the ramped current pulse is only slightly less efficient in stimulating the heart than the constant current pulse. The use of a ramped current waveform is believed to provide a non-invasive pacemaker which can generally be tolerated for longer periods of time than a constant pulse generator.

The inventors have found that the response to external non-invasive pacing is highly individualized. Although generally muscle stimulation is reduced by extending the ramp time of the pacing pulse, the effect of such an extension varies from patient to patient. Further, the length of time a patient can tolerate external pacing is also related to the pain induced by the pacemaker. Pain also varies in a complex fashion with ramp time and differs from patient to patient. Therefore, the optimum ramp time varies from patient to patient. In the pacemaker of the present invention the ramp time preferably is adjustable to allow the pacing therapy to be optimized for each patient. In conformity with a preferred further development the pacemaker allows adjustment of the ramp time and maximum pulse current amplitude to provide the most comfortable waveform consistent with reliably pacing the heart.

An exemplary embodiment of the invention will be described with reference to the accompanying drawings, in which:

Fig. 1 illustrates a typical prior art constant current output waveform.

Fig. 2 illustrates a ramped current waveform according to the present invention.

Fig. 3 including Figs 3a, 3b, 3c and 3d illustrates the schematic of a pacemaker according to the present invention which generates the pacing pulse of Fig. 2.

Fig. 4 is the schematic of the power supply for a pacemaker according to the present invention.

Fig. 1 shows a typical prior art constant current waveform of the sort used with present day external non-invasive cardiac pacemakers. Typically, the pulse is 20 to 40ms in duration. The rise time 10 and the fall time 14 of the pulse are typically on the order of 1 to 2ms. The pulse rapidly attains its predetermined maximum current level 12 rapidly and substantially maintains that current level throughout the duration of the pulse.

Fig. 2 illustrates the output current pulse of a pacemaker according to the present invention. The overall duration of the pulse should be at least 20ms, and preferably at least about 40ms.

The current pulse of Fig. 2 differs from the prior art in that the pulse, instead of rising rapidly to its maximum current level 20, employs an extended, gradual rise. The rise time 16 of the ramped waveform if fixed should be a substantial portion of the duration of the waveform and, preferably, at least half of the duration of the waveform. The total duration of the waveform if fixed should be at least about 40ms.

- 5 Generally, the maximum reduction in muscle stimulation occurs with a fully ramped waveform, e.g. 40ms duration and approximately a 40ms rise time. However, pacing efficiency starts decreasing more significantly with rise times in excess of 30ms. If rise time is also fixed, a rise time of between 20 and 40ms is believed advisable with 30-40ms preferable.

- 10 After reaching the maximum current level 20, the output pulse continues at this current level for a short period of time 18, and then rapidly returns to 0 over a short fall time 21 which may be 1 or 2ms. The time period 18 during which the pulse is at its maximum current level if fixed is preferably 0-10ms. Based on experimentation performed by the inventors, it appears that extending the fall time 21 of the ramped waveform generally provides no particularly additional benefits, but may be desirable in some patients.

Fig. 3 illustrates the interconnection between the schematic diagrams of Figs. 3a, 3b, 3c, and 3d.

- 15 Fig. 3a illustrates the circuitry associated with the sensing of the patient's heartbeats. EKG electrodes 100 and 101 are coupled via resistors 102 and 103 and coupling capacitors 148 and 149 to the input of amplifier 109. Diodes 104 and 105 serve to protect amplifier 109 from the application of high voltages to electrodes 100 and 101. Diodes 150 and 151 limit the input voltages to amplifier 109 to one diode drop. Coupling capacitors 148 and 149 allow electrodes 100 and 101 to also be used to pace the heart. Amplifier 20 109, in conjunction with associated resistors 106, 107, 108, 110, 111, 112, 113, and 114, and capacitor 116, provides the initial amplification of the EKG signal. The DC offset of amplifier 109 is determined by resistors 106, 107, 108, 145 and 114, and adjusted by resistor 107. The common mode rejection capability of amplifier 109 is determined by resistors 112 and 113 in conjunction with capacitor 116. Adjustment is accomplished by resistor 113. Overall, amplifier 109 provides approximately a 2X gain.

- 25 The output of amplifier 109 is passed via resistor 117 and capacitor 118 to amplifier 121. Resistor 117 and capacitor 118 act as a low pass filter, allowing frequencies of approximately 70Hz and below to pass through to amplifier 121. Amplifier 121, in conjunction with associated resistors 115 and 120, functions as a noninverting amplifier, with a gain of approximately 50X. The output of amplifier 121 passes via resistor 122 and capacitor 123 to the input of amplifier 127. Amplifier 127, in conjunction with associated resistors 125 30 and 126 and capacitors 123 and 124, act as a band pass amplifier with a gain of approximately 4X to 8X. The band pass defined by the circuitry extends from approximately .5Hz to approximately 40Hz.

- The output of amplifier 127 is passed through to amplifiers 136 and 137 which act as reference comparators, comparing the negative and positive amplitude of the output of amplifier 127 to positive and negative reference voltages. Amplifier 131, in conjunction with associated resistors 129 and 130, provides 35 comparing amplifier 137 with a positive reference voltage adjustable from .5mv to approximately 10mv. R128 and R129 serve as internal high end and low end calibration resistors. Resistor 130 serves as the sensitivity control for the pacemaker. The output of amplifier 131 is also passed via resistor 132 to amplifier 133 which, in conjunction with associated resistors 134 and 135, sets the negative reference voltage provided to comparing amplifier 136. Adjustable resistor 135 provides an internal calibration adjustment to 40 minimize the disparity between the positive and negative reference signals. If either the positive or negative going portion of the signal provided by amplifier 127 exceeds the reference value, an output signal is generated indicative of a sensed ventricular contraction and, in conjunction with resistors 138 and 139 and diodes 140 and 141, is passed through to reset the timing circuitry, discussed below. Resistors 142 and 143 serve as pull-up resistors. Capacitors 152 and 153 serve provide a cleaner reset signal from the comparing 45 amplifiers 136 and 137. Switch 146 is coupled to adjustable resistor 130, which serves as the sensitivity control. When switch 146 is open, no sensing of heart activity is accomplished, and the pacemaker operates in an asynchronous mode. Resistor 144 ties line 147 to ground in the event that asynchronous pacing is selected.

- Fig. 3b shows the circuitry associated with the basic timing of pacing pulses. The retriggerable one- 50 shots 200 and 202, along with their associated capacitors 204 and 206 and resistors 208, 210 and 212 serve to determine the basic pacing rate. Variable resistor 208 serves as the pacemaker's rate control. The timing interval determined by one-shot 200 should be variable at least between about 1200ms and 500ms, but pacing intervals as short as 300ms may be desirable for some uses. The negative edge of the signal from comparator amplifiers 136 and 137 on line 147 reinitiates the timing period of one-shot 200, which 55 determines the basic pulse interval and sets the Q output of one-shot 200 high. On the expiration of the timing interval determined by capacitor 204 and resistors 208 and 210, the Q output of one-shot 200 goes low. Oscillator 241, in conjunction with associated capacitor 245 and resistor 243, generates a 30KHz squarewave on line 160. When the Q output of one-shot 200 goes low, or on the positive going transition on

line 160 thereafter, one-shot 202 is initiated. One-shot 202 then times out a 5ms interval determined by resistor 212 and capacitor 206, generating a 5ms low logic level pulse on its \overline{Q} output which retriggers one-shot 200. As such, the pair of one-shots form a retriggerable, free running oscillator. The corresponding 5ms high pulse on the Q output of one-shot 202 triggers non-retriggerable one-shot 214 driving its Q output high.

One-shot 214, in conjunction with associated capacitor 216 and resistor 218, determines a 100ms interval during which the pacing pulse is generated. The low logic level pulse at the \overline{Q} output of one-shot 214 serves to trigger the generation of a pacing pulse. The positive going transition of the high logic level output pulse from the Q output of one-shot 214 is used to trigger a second non-retriggerable one-shot 220, which serves as the rate limit for the pacemaker. The high logic level pulse on the Q output of one-shot 214 also activates transistor 215 to turn on LED 219 via resistor 217. LED 219 serves as a visual indicator of the generation of a pacing pulse.

One-shot 220, in conjunction with associated capacitor 222 and resistor 224 generates a 260ms high logic level pulse on the Q output of one-shot 220 which prevents further triggering of one-shot 214. This arrangement prevents any malfunction of the rate control one-shots 200 and 202 from causing the generation of pacing pulses in excess of about 230 beats per minute. Adjustment of the value of resistor 224 would allow for selection of the higher or lower rate limits.

The high logic level pulse on the Q output of one-shot 214 also serves to set the output of OR gate 221 high, which in turn couples line 147 to ground via digital switch 232. This prevents subsequent sensed contractions of the heart from restarting the timing period for at least the 260ms period determined by one-shot 214, thus provided a post-pacing blanking period.

Each time retriggerable one shot 200 is triggered by the sensing of a signal from comparator amplifiers 136 and 137 (Fig. 3a), non-retriggerable one-shot 234 is also triggered. One shot 234, in conjunction with capacitor 236 and resistor 238 generates a 100ms high logic level signal on its Q output. This turns on transistor 240, allowing current to flow through LED 242 and resistor 244. LED 242 indicates the sensing of a heartbeat. The visual indication of heartbeat sensing assists in adjustment of the sensitivity control, resistor 130 (Fig. 3a) to an appropriate level to reliably sense the heartbeat.

The Q output of one-shot 234 going high also sets the output of OR gate 221 high, tying line 147 low via digital switch 232 and associated resistor 233. Thus, one-shot 234 also provides a 100ms post-sensing blanking interval. Blanking functions can be disabled via switch 235.

Fig. 3c illustrates the high voltage oscillator and the oscillator control circuitry. Multivibrator 246 in conjunction with timing resistor 248 and timing capacitor 250 is configured to provide 133KHz square wave signals on its Q and \overline{Q} outputs, 90 degrees out of phase with one another. Amplifiers 252 and 254 are used as FET drivers to turn FET's 256 and 258, respectively, on and off in response to the signals generated on the Q and \overline{Q} of oscillator 246. FET's 256 and 258 provide out of phase 12 volt, 133KHz signals to transformer 260. Transformer 260 steps up the ± 12 volt signals to a ± 200 volt signal, which is subsequently rectified by the diode bridge including diodes 270, 272 274 and 276 and stores in capacitor 278.

The voltage stored in capacitor 278 is divided by a factor of 100 by resistors 280 and 282 which form a voltage divider. The divided voltage is compared by differential amplifier 284 to a 2 volt reference signal provided by resistor 283 and diode 285. When the voltage stored on capacitor 278 reaches 200 volts, oscillator 246 is turned off via differential amplifier 286, which generates a reset signal on line 288. The reset signal on line 288 sets the output of exclusive OR gate 290 low so that FET 258 will also be turned off when the oscillator is held on reset. This feature avoids unnecessary battery drain.

Differential amplifier 286, in conjunction with associated resistors 292, 294, 296 and 298 also serves as a comparator. Differential amplifier 286 is configured to also generate a reset signal for oscillator 246 in response to a high logic level signal on line 300, which is coupled to the output circuit and is high during generation of a pacing pulse. This prevents any signals from oscillator 246 from being superimposed on the pacing pulse. Capacitor 302 and resistor 304 serve to filter high frequency switching artifacts from the reset signal.

During generation of the pacing pulse, capacitor 278 is discharged, and passed through the output circuit. Switch 306 disconnects capacitor 278 from the output circuit. Switch 308 allows capacitor 278 to be discharged via resistor 310, and is provided to facilitate servicing the pacer.

Fig. 3d illustrates the pulse generator of the 30 pacemaker, including circuitry for generating the ramped output waveform. One-shot 315, in conjunction with associated resistor 317 and capacitor 319, determines the pulse width of the pacemaker. Resistor 317 serves as an adjustment for the pulse width, which may be adjustable between 0 and 40ms, or at least between 20 and 40ms.

Differential amplifier 312 and associated resistors 314, 316, 318 and capacitor 320 generate the ramp waveform. The 100ms square wave generated on the \overline{Q} output of one-shot 214 (Fig. 3B) triggers one-shot

315.

The negative going logic pulse generated on the \overline{Q} output of one-shot 315 is integrated by differential amplifier 312 and its associated circuitry, including resistors 316 and 318 and capacitor 320. Variable resistor 314 provides an adjustable rise time ramp. Variable resistor 321, in conjunction with diode 323, adjust the fall time of the output pulse.

The ramped waveform provided by amplifier 312 is provided to amplifier 326 via resistor 324 and diode 322. Amplifier 326 is configured as a constant current amplifier. The output of amplifier 326 is used to control FET 328, which regulates the current flow through the patient. As such, amplifier 326 in conjunction with FET 328 provides a ramped current waveform, having ramp determined by the output of amplifier 312. Resistor 330 is used to regulate the output current, and resistors 332, 334 and capacitor 336 are used to filter out any transients generated by amplifier 326. Resistor 338 allows amplifier 326 to sample the current flowing through FET 328 and thus regulate the current.

FET 340 is used to ground patient electrode 342 between pacing pulses. FET 340 is turned off by amplifier 344 during pacing pulses. Resistor 346 and diode 348 ensure that FET 340 is turned on slowly following the pacing pulse to avoid clipping the pacing pulse. The output of comparing amplifier 344 is also provided to amplifier 286 (Fig. 3c) and is used to disable oscillator 246 (Fig. 3c) during the pacing pulse, as discussed above. Resistor 381 acts as a pull-up resistor. Capacitor 373 is used for common mode rejection.

FET 352 and associated transistor 354 and resistors 356 and 358 are used to connect the output of capacitor 278 (Fig. 3c) to patient electrode 342 during a pacing pulse. FET 352 is turned on by FET 360, which is controlled by amplifier 362. Comparing amplifier 362 turns on transistor 360 in response to the ramp output of amplifier 312 during the pacing pulse. Resistor 375 acts as a pull-up resistor. Capacitor 371 is used for common mode rejection. Diode 364 and resistor 366 ensure that FET 360 is turned off slowly. Transistor 354 is turned on whenever the current through FET 352 exceeds 200 milliamps, and serves to limit the current to 200 milliamps in case of component failure elsewhere in the circuit.

The output circuitry of Fig. 3d will require modification in those jurisdictions in which A-C coupled output stages are required by governmental regulation. Such modifications are believed to be well within the skill in the art.

Fig. 4 illustrates the power and reference circuits used in the pacemaker. The battery provides a 12 volt power supply via diode 377 and resistor 379. Voltage regulator 368 and associated capacitors 370, 372, 374, and diodes 376 and 378 convert the 12 volt signal from the battery to the positive 5 volt signal used to drive the digital circuitry in the pacemaker. Capacitors 370, 372, 374 and diodes 376 and 378 provide filtering and voltage limiting.

Voltage regulator 380 and associated resistors 382, 384, 386 and 388 provide a positive 5 volt signal used by the linear circuitry in Figs. 3a through 3d. Variable resistor 386 is used to calibrate the positive 5 volt signal.

The 12 volt signal provided by the battery is applied to voltage inverter 390 via LED 392, which acts as an indicator that the pacemaker is turned on. Voltage inverter 390, in conjunction with capacitors 394, 396 and 298, inverts the 12 volt signal from the battery and applies it to voltage regulator 400 via diode 402. Voltage regulator 400, in conjunction with resistors 404, 406, 408 and 410 and capacitors 412, 414 and 416, provides a negative 5 volt power signal to the linear circuitry illustrated in figures 3a-3d. Variable resistor 406 is used for calibration of the positive 5 volt signal.

The design of the pacemaker allows the pacing pulse configuration to be adjusted to each patient. Generally, pain and muscle stimulation increase with increased pulse amplitude, but can often be reduced by extending the ramp time of the output pulse. Extending the duration of the output pulse generally allows use of lower pacing amplitudes. Generally, adjustment of the device should involve starting with a fully ramped waveform with a rise time of at least about 40ms and adjusting the amplitude control resistor 330 to the lowest maximum current level consistent with reliable capture of the heart, as indicated by an EKG monitor. The ramp time and amplitude may then be adjusted to determine whether a lesser ramp time may allow further reduction in the pulse amplitude required to reliably pace the heart. Adjustment of the amplitude and ramp times should continue until the patient indicates his maximum comfort level consistent with reliable pacing. After pacing for a period of time, the pacemaker may be readjusted if necessary. It is anticipated that some patients' ability to tolerate pacing for extended periods of time may be improved merely by regularly varying the pulse waveform.

COMPONENT LISTINGAmplifiersVoltage Regulators

109	LH0036C	368	LM2931A
121,127,131,133	LM224	380	1CL7663
136,137,362,344	LM393	390	1CL7660
284	TLC27L4	400	1CL7664
326	LM10C		
312,286	TLC27M4		

Logic Gates

290	CD4030
252,256	CD4050
221	CD4075B

Oscillators

241,246	CD4047
---------	--------

			<u>Diodes</u>	
	<u>One-Shots</u>		104, 105, 150, 151	1N4007
5	200, 202	74HC423	140, 141	1N4148
	214, 220, 234	74HC221	323, 364, 348, 402	1N5711
	315	CD4098	322	1N750
10			376	1N5243
	<u>Digital Switch</u>		378	1N5235
	232	CD4066	270, 272, 274, 377	1N4007
			264, 266, 276	1N4007
15	<u>Transistors</u>		262, 268	1N974
	240, 215	2N2222	285	LM113
	354	2N2907		
20	352	MTP2P45	<u>LED's</u>	
	360	VN2406L	392	MLED600
	340, 328	1RFF321	219	LDH5191
25	256, 258	1RFD111	242	LDY5391
	<u>Resistors (Ω)</u>		<u>Capacitors (F)</u>	
30	106, 108, 132	100K	116	.01u
	107, 130, 135	100K Adj.	123, 372, 416	.047u
	145	33K	124	.0047u
	102, 103, 366	20K	204	15u
35	112	52.3K	206	1000p
	114	4.02K	216, 222, 118, 236	.1u
	120	250K Adj.	245	120p
40	117	24.3K	250, 394	100p
	110, 111, 125	1M	278	150u
	122	88.7K	414, 302, 374	100u
45	128	500K Adj.	319	1.0u
	129	10K Adj.	320	.27u
	134	56K	336	10pf
	142, 143, 139	100K	370	2200u
50	138	510K	412, 398, 396	68u
	126, 144, 238	1M	148, 149	.22u
	244	240	152, 153, 371, 373	.001u
55	233, 218	1M		

Resistors (Ω) Cont.

5	208,330	100K Adj.
	210	21K
	212,294,375,381	5K
	224	2.6M
10	217	500
	243	57.6K
	248	75K
15	310	700
	282	220K
	280	41.2M
20	292,296,298	50K
	283,304,334	20K
	317	500K Adj.
25	314,321	200K Adj.
	316,318,332	100K
	324	210K
	338	10
30	346	240K
	347,357	1M
	356	51K
35	358	2
	410,379,382	20
	384,404	1.5M
40	388,408	470K
	386,406	200K Adj.
	367	100K
45	369	100

Claims

1. A non-invasive cardiac pacemaker, comprising:
 first and second skin contacting electrode means for electrically coupling to a patient;
 first timing means for repetitively timing an escape interval; and
 pulse generator means, coupled to and responsive to said first timing means and coupled to said first and
 second skin contacting electrode means, said pulse generator means generating pacing pulses for
 application to said patient via said first and second electrode means, said pacing pulses extending for a

duration of at least about 20ms, said pacing pulses having a current amplitude which rises gradually from 0 to a maximum current amplitude over a predetermined rise time of at least about 20ms, said predetermined rise time extending over at least about one-half of the duration of said pacing pulse.

2. A pacemaker according to claim 1 wherein said pulse generator means pacing pulses of at least about 40ms in duration, and in which the rise time of said pacing pulses extends for at least about 30ms.

3. A pacemaker according to claim 2 wherein the rise time of said pacing pulses extends for about 40ms.

4. A pacemaker according to claim 1 or claim 2 or claim 3 wherein the current amplitude of said pacing pulses pulse from said maximum current level to 0 in about 2ms or less at the expiration of said pacing pulse.

5. A pacemaker according to claim 1 wherein said pulse generator means further includes means for adjusting the rise time of said pacemaker pulses.

6. A pacemaker according to claim 5 wherein said pulse generator means generates pacing pulses having a duration of about 40ms and wherein said rise time of said pacemaker pulses is adjustable at least between 30 and 40ms.

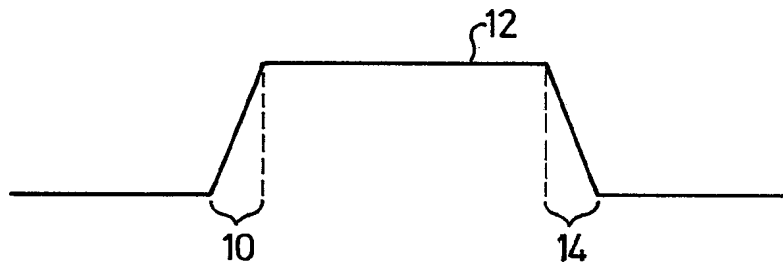


FIG. 1 (PRIOR ART)

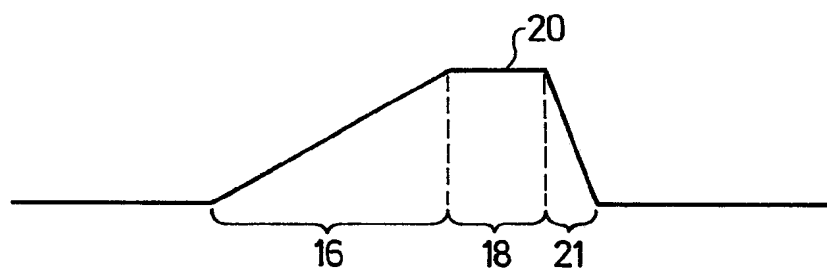
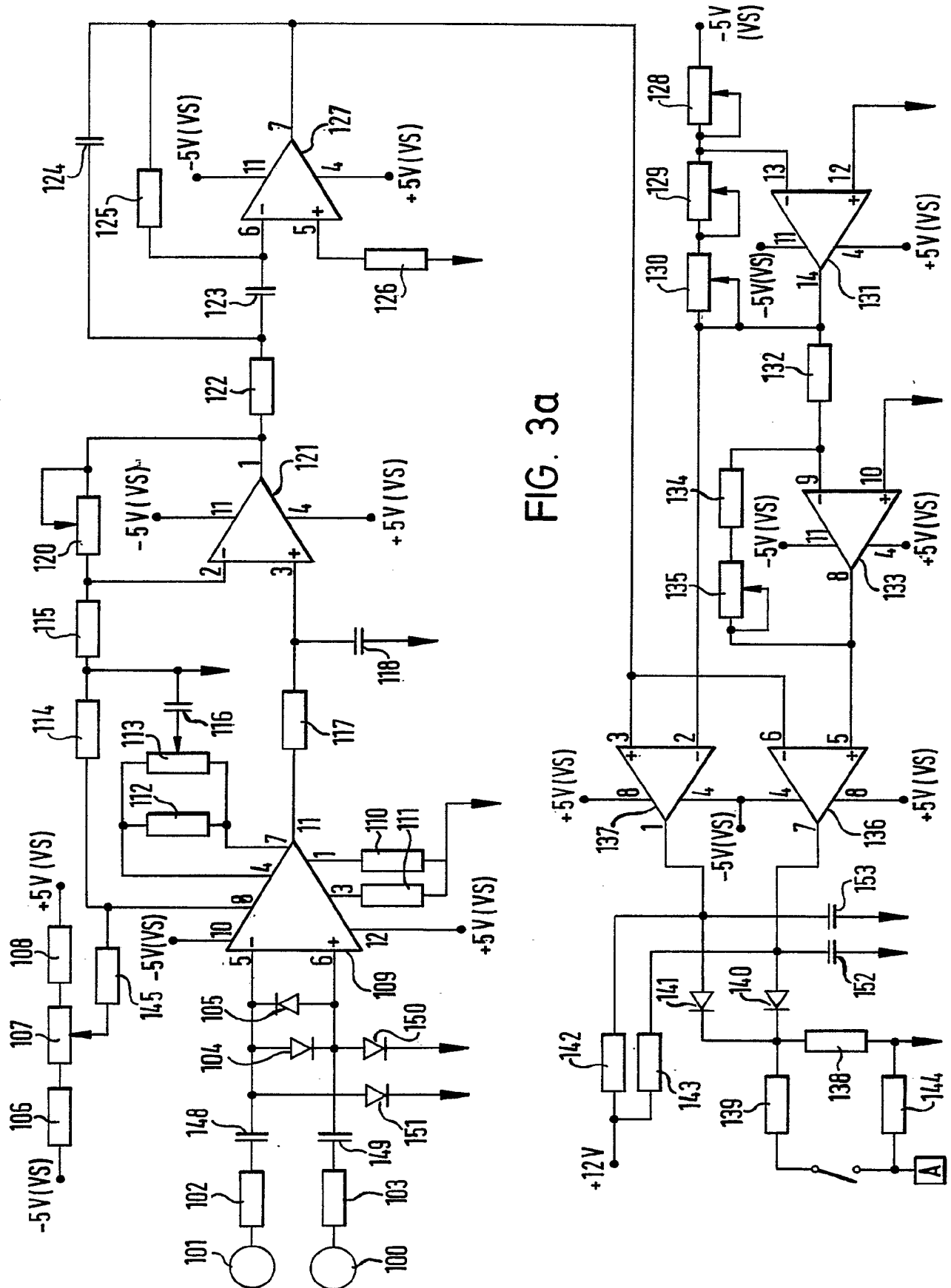


FIG. 2

3a	3c
3b	3d

FIG. 3



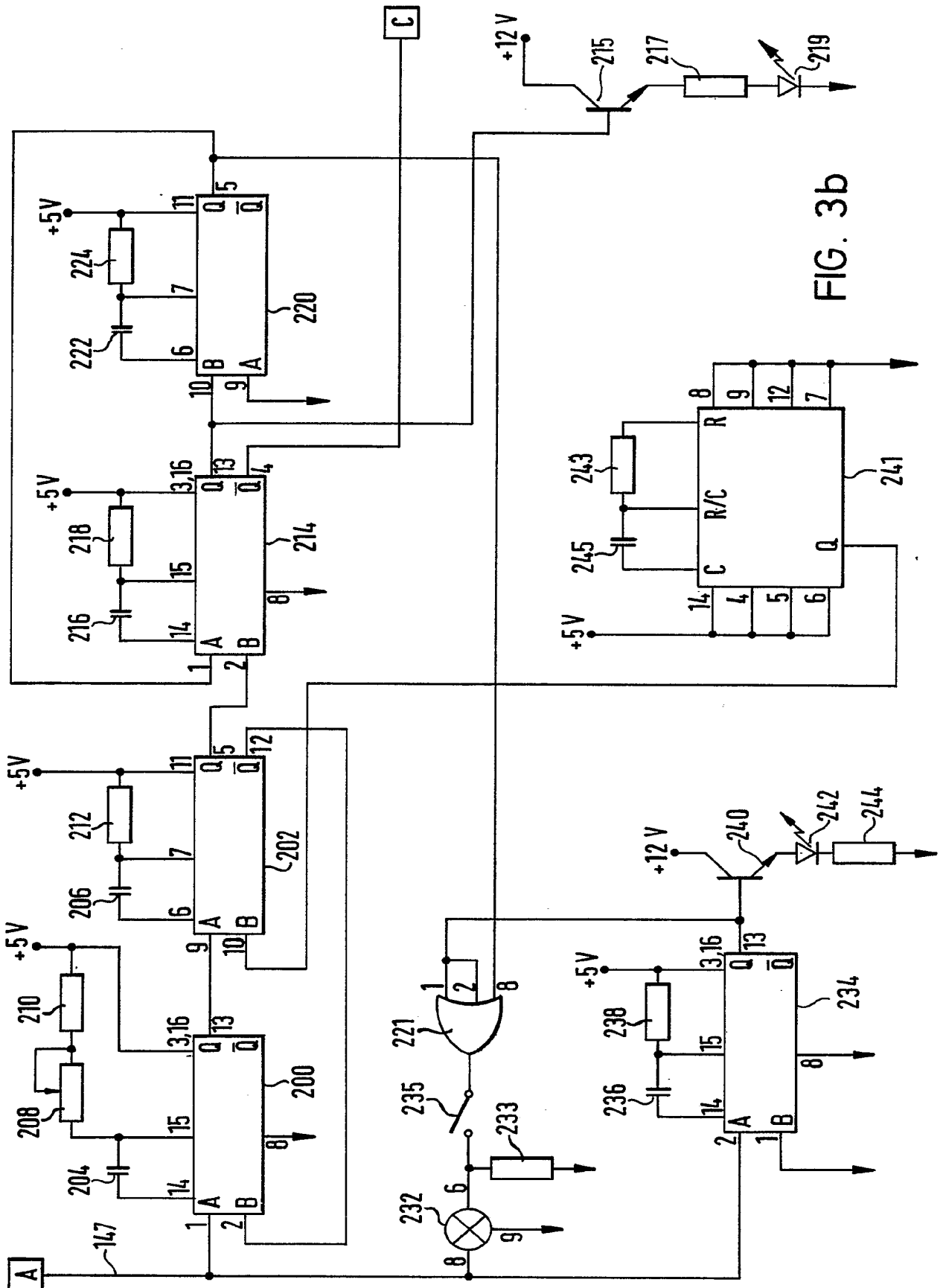
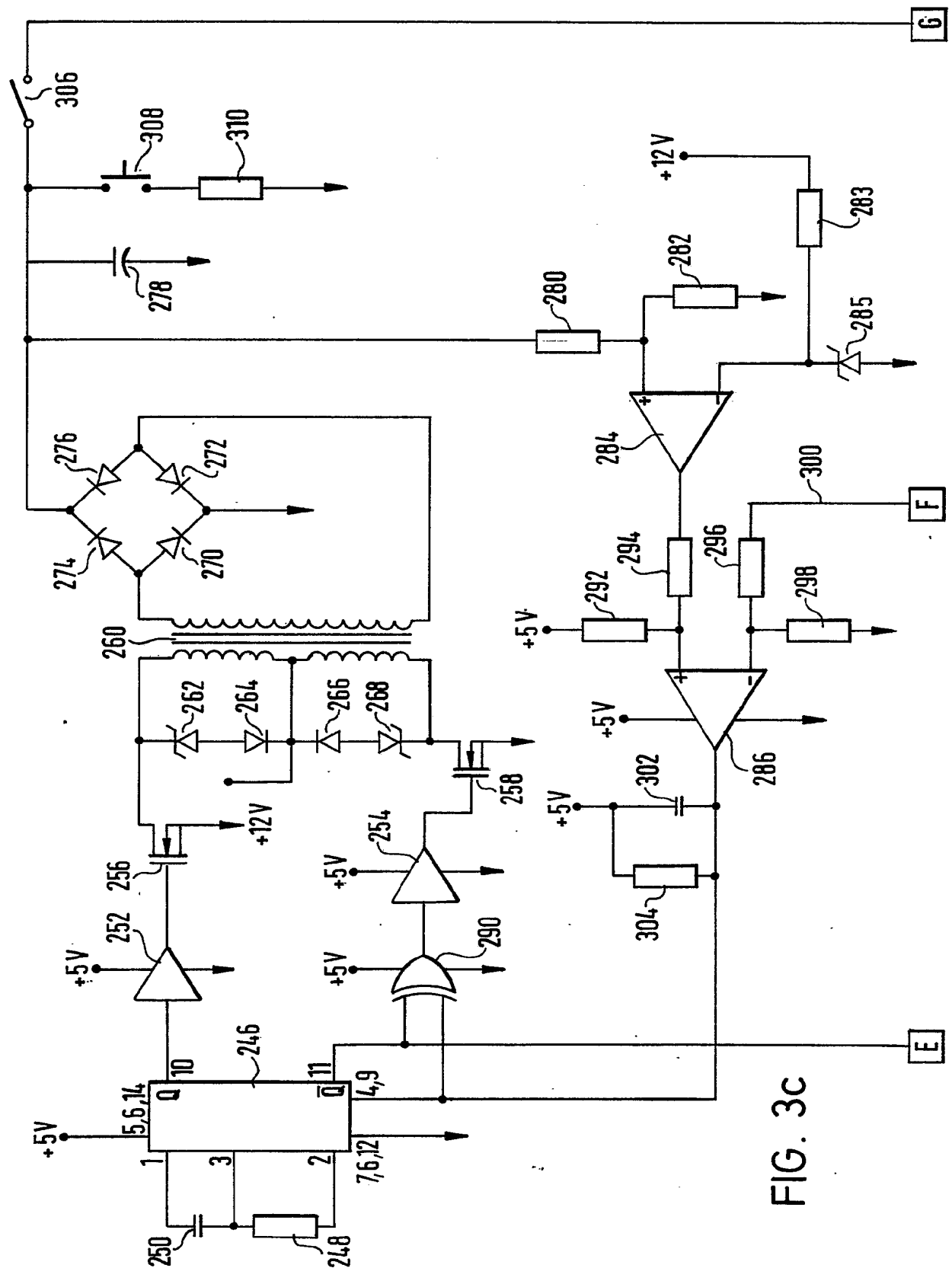


FIG. 3b



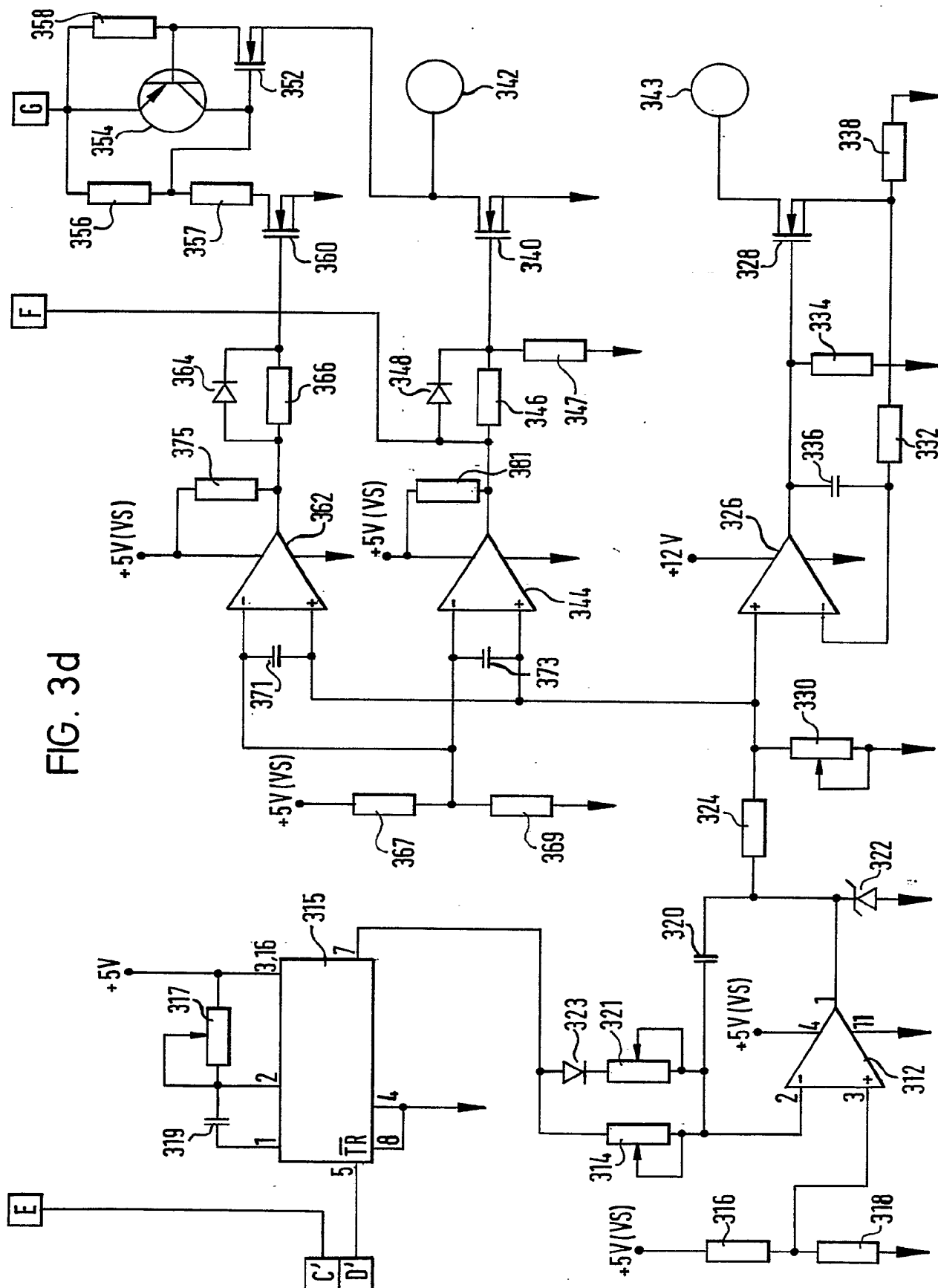
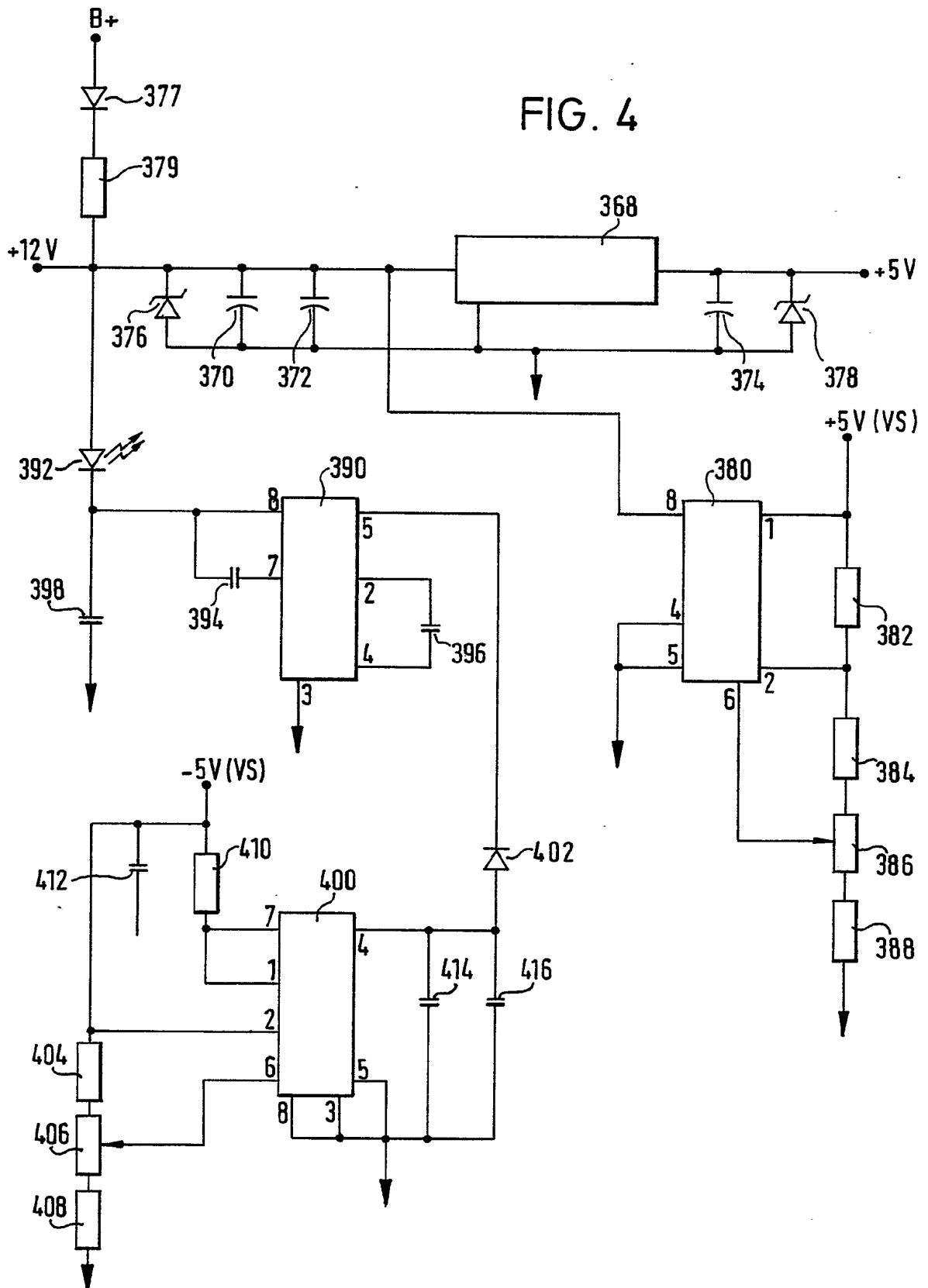


FIG. 4





European Patent
Office

EUROPEAN SEARCH REPORT

Application Number

EP 88 11 7771

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.4)
Y,D	US-A-4 349 030 (BELGARD) * Column 1, lines 23-46; column 2, lines 64-66; column 4, lines 3-5; column 4, lines 8-9 *	1,4,5	A 61 N 1/362
A	---	2,3,6	
Y	US-A-4 535 777 (CASTEL) * Column 1, line 65 - column 2, line 42; column 4, lines 37-63 *	1,4,5	
A	---	2,3,6	
A,P	US-A-4 735 206 (HEWSON) * Column 4, lines 56-66; figure 5 * -----	1,2,3,6	
			TECHNICAL FIELDS SEARCHED (Int. Cl.4)
			A 61 N
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 18-01-1989	Examiner LEMERCIER D.L.L.
CATEGORY OF CITED DOCUMENTS X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document			